

# Enrocin 10% Oral 100 mg/ml Roztwór do podania w wodzie do picia

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

Enrocin 10% Oral 100 mg/ml Roztwór do podania w wodzie do picia

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**Active substance:**

Enrofloxacin

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**Target species:**

Pigeon

Chicken (hen)

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**Route of administration:**

In drinking water use

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## Product details

**Active substance and strength:**

Enrofloxacin

100.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for use in drinking water

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**Withdrawal period by route of administration:****In drinking water use:**

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**Pigeon**

- All relevant tissues. no withdrawal period

The product is not approved for use in birds producing eggs for human consumption. Do not use in young birds reared for laying within 14 days before the onset of the laying period. Do not use in pigeons intended for consumption.

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**Chicken (hen)**

- Meat and offal. 7 day

The product is not approved for use in birds producing eggs for human consumption. Do not use in young birds reared for laying within 14 days before the onset of the laying period. Do not use in pigeons intended for consumption.

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01MA90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Poland

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**Available in:**

Poland

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**Package description:**

Available only in Polish

Available only in Polish

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Well-established use application (Article 13a of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

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**Marketing authorisation date:**

27/03/2001

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**Manufacturing sites for batch release:**

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

1148

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**Date of authorisation status change:**

27/03/2001

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Package Leaflet

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## Labelling

This document does not exist in this language (English). You can find it in another language below.

## Summary of Product Characteristics

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